SUPPLEMENTAL INFORMATION DISCLOSURE STATEMENT LIST

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Complete if Known	
Application Number	10/579,149
Filing Date	January 19, 2007
First Named Inventor	Kortney L. Klinkel
Group Art Unit	1611
Examiner Name	Joseph S. Kudla

U.S. PATENT DOCUMENTS

Examiner's Initials	Cite No.	Document No.	Date	Name	Class	Subclass	Filing Date (if appropriate)
	B1	3,279,918	10-18-1966	Gevaert Photo-Production N.V.	90	1	
	B2	3,297,710	1-10-1967	E.I. du Pont de Nemours & Co.	260	309	
	B3	4,089,747	5-16-1978	Eastman Kodak Co.	435	10	
	B4	4,423,046	12-27-1983	Sterling Drug Inc.	514	228.8	
	B5	4,466,976	8-21-1984	Schering Aktiengesellschacft	514	397	
	B6	4,585,771	4-29-1986	Schering Aktiengesellschacft	514	220	
	B7	4,705,855	11-10-1987	Rottapharm SpA	544	370	
	B8	4,721,670	1-26-1988	Fuji Photo Film Co.	435	28	
	B9	4,902,705	2-20-1990	UBE Industries, Ltd. et al.	514	397	
	B10	4,970,226	11-13-1990	Harbor Branch Oceanographic Institution, Inc.	514	397	
	B11	5,024,935	6-18-1991	Eastman Kodak Co.	435	7.1	
	B12	5,047,318	9-10-1991	Eastman Kodak Co.	435	5	
	B13	5,496,702	3-5-1996	Johnson & Johnson Chemical Diagnostics, Inc.	435	7.9	
	B14	5,514,550	5-7-1996	Johnson & Johnson Chemical Diagnostics, Inc.	435	6	
	B15	5,656,644	8-12-1997	SmithKline Beecham Corp.	514	341	
	B16	5,686,455	11-11-1997	SmithKline Beecham Corp.	514	256	
	B17	5,693,589	12-2-1997	Eastman Kodak Co.	503	227	
	B18	5,753,687	5-19-1998	Ontogen Corp.	514	396	
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	B20	5,945,418	8-31-1999	Vertex Pharmaceuticals, Incorp.	514	248	
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	B24	6,268,370	7-31-2001	SmithKline Beecham Corp.	514	256	
	B25	6,521,655	2-18-2003	Ortho-McNeill Pharmaceuticals, Inc.	514	397	
	B26	2007/0123553	5-31-2007	Lorus Therapeutics Inc.	514	285	
	B27	2004/0127527	7-1-2004	Mitsuya et al.	514	365	
	B28	2004/0176601	9-9-2004	Goulet et al.	546	46	
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	B31	EP 165588	12-27-1985	Fuji Photo Film Co. Ltd.	
	B32	JP 2002-275458	9-25-2002	Fukuoka Prefecture	Abstract only
	B33	JP 2002-364578	12-18-2002	Hitachi Ltd.	Abstract only
	B34	WO 1995/03297	2-2-1995	SmithKline Beecham Corp.	
	B35	WO 1996/18626	6-20-1996	F. Hoffman-La Roche AG Harmon	
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	B37	WO 1999/01128	1-14-1999	Neurogen Corp.	
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	B39	WO 2003/004023	1-16-2003	Schering Aktiengesellschacft	
	B40	WO 1993/014081	7-22-1993	SmithKline Beecham Corp.	
	B41	WO 1998/027065	6-25-1998	Ontogen Corp.	
	B42	WO 2000/078761	12-28-2000	Sepracor Inc.	
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	B45	WO 2005/047266	11-15-2004	Lorus Therapeutics Inc.	

NON-PATENT DOCUMENTS		
Examiner's Initials	Cite No.	Non-Patent Citations (include Author, Title, Publisher, Relevant Pages, Date and Place of Publication)
	B46	Abdel-Meguid *et al.*, "An orally bioavailable HIV-1 protease inhibitor containing an imidazole-derived peptide bond replacement: Crystallographic and pharmacokinetic analysis," *Chemistry*, 1994, 33:11671-11677
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	B48	Antolini *et al.*, "Analogues of 4,5-bis(3,5-dichlorophenyl)-2-trifluoromethyl-1H-imidazole as potential antibacterial agents," *Bioorg. Med. Chem. Lett.*, 1999, 9:1023-1028
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	B50	Bhaduri *et al.*, "Potential Antifertility Agents. Synthesis of 2,4,5-Substituted Imidazoles", Central Drug Res. Inst. Lucknow, India, *Indian J. Chem.*, 1966, 4(9):419-420
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	B58	Demirayak *et al.*, "Synthesis of Certain Derivatives of Ethyl α-[(phenanthro[9,10-d]imidazol-2-yl)phenoxy]alkanoate", *Acta Pharmaceutica Turcica*, 1989, 31(1):19-25
	B59	Downey *et al.*, "Degradation of DNA by 1,-10-phenanthroline," *Biochem Biophys Res Commun*, 1980, 93(1):264-70
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	B66	Kimura *et al.*, "*Preparation of 4-(4,5-diphenyl-1H-imidazol-2-yl)benzaldehyde and its Practical Synthetic Use in the Synthesis of Unsymmetrically Substituted Imidazoles*", Department of Chemistry, Okayama University, Okayama Japan, ITE Letters on Batteries, New Technologies & Medicine, 2002, 3(1), pp. 30-34
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